Transport Spectroscopy of a Kondo quantum dot coupled to a finite size grain.

Pascal Simon¹, Julien Salomez¹ and Denis Feinberg²

¹Laboratoire de Physique et Modélisation des Milieux Condensés,

CNRS et Université Joseph Fourier, 38042 Grenoble, France

² Laboratoire d'Etudes des Propriétés Electroniques des Solides, CNRS,
associated with Université Joseph Fourier, BP 166, 38042 Grenoble, France

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We analyse a simple setup in which a quantum dot is strongly connected to a metallic grain or finite size wire and weakly connected to two normal leads. The Kondo screening cloud essentially develops in the strongly coupled grain whereas the two weakly connected reservoirs can be used as transport probes. Since the transport channels and the screening channels are almost decoupled, such a setup allows an easier access to the measure of finite-size Kondo effects.

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I. INTRODUCTION

One of the most remarkable achievements of recent progress in nanoelectronics has been the observation of the Kondo effect in a single semi-conductor quantum dot. 1,2,3 When the number of electrons in the dot is odd, it can behave as an S=1/2 magnetic impurity interacting via magnetic exchange with the conduction electrons. One of the main signatures of the Kondo effect is a zerobias anomaly and the conductance reaching the unitary limit $2e^2/h$ at low enough temperature $T < T_K^0$. T_K^0 stands for the Kondo temperature and is the main energy scale of the problem. At low temperature, the impurity spin is screened and forms a singlet with a conduction electron belonging to a very extended many-body wavefunction known as the Kondo screening cloud. The size of this screening cloud may be evaluated as $\xi_K^0 \approx \hbar v_F/T_K^0$ where v_F is the Fermi velocity. In a quantum dot, the typical Kondo temperature is of order 1 K which leads to $\xi_K^0 \approx 1$ micron in semiconducting heterostructures. Finite size effects (FSE) related to the actual extent of this length scale have been predicted recently in different geometries: an impurity embedded in a finite size box,⁴ a quantum dot embedded in a ring threaded by a magnetic flux, 5,6,7,8 and a quantum dot embedded between two open finite size wires (OFSW) (by open we mean connected to at least one external infinite lead). 9,10 In the ring geometry, it was shown that the persistent current induced by a magnetic flux is particularly sensitive to screening cloud effects and is drastically reduced when the circumference of the ring becomes smaller than ξ_K^0 . In the wire geometry, a signature of the finite size extension of the Kondo cloud was found in the temperature dependence of the conductance through the whole system.^{9,10} To be more precise, in a one-dimensional geometry where the finite size l is associated to a level spacing Δ , the Kondo cloud fully develops if $\xi_K^0 \ll l$, a condition equivalent to $T_K^0 \gg \Delta$. On the contrary, FSE effects appear if $\xi_K^0 > l$ or $T_K^0 < \Delta$.

Nevertheless, in such a two-terminal geometry, the screening of the artificial spin impurity is done in the OFSWs which are also used to probe transport properties through the whole system. This brings at least two main drawbacks: first, the analysis of FSE relies on the independent control of the two wire gate voltages and also a rather symmetric geometry. This is difficult to achieve experimentally. In order to remedy these drawbacks, we propose and study here a simpler setup in which the screening of the impurity occurs mainly in one larger quantum dot or metallic grain¹¹ or OFSW and the transport is analyzed by help of one or two weakly coupled leads. In practice, a lead weakly coupled to the dot by a tunnel junction allows a spectroscopic analysis of the dot local density of states (LDOS) in a way very similar to a STM tip. The rather general geometry we study is depicted in Fig. 1. We note that this geometry has also been proposed by Oreg and Goldhaber-Gordon¹² to look for signatures of the two-channel Kondo fixed point or by Craig et al. 13 to analyze two quantum dots coupled to a common larger quantum dots and interacting via the RKKY interaction. In the former case, the key ingredient is the Coulomb interaction of the grain whereas in the latter, the grain is largely open and used simply as a metallic reservoir mediating both the Kondo and RKKY interactions. Here we are interested in the case where finite-size effects in the larger quantum dot or metallic grain do matter whereas in the aforementioned experiments the level spacing was among the smallest scales.

The plan of the paper is the following: in section 2, we present the model Hamiltonian and derive how the FSE renormalizes the Kondo temperature in our geometry. In section 3, we show how FSE affect the transport properties of the quantum dot and perform a detailed spectroscopic analysis. The effect of a finite Coulomb energy in the grain is also discussed. Finally section 4 summarizes our results.

II. PRESENTATION OF THE MODEL

A. Model Hamiltonian

The geometry we analyze is depicted in Fig. 1. In this section we assume that the large dot is connected to a

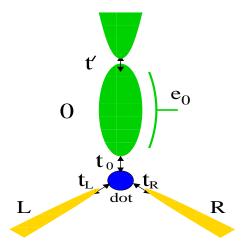


FIG. 1: Schematic representation of the device under analysis. When the grain Coulomb blockade energy is not neglected, we assume that the grain potential can be controlled by a voltage gate e_0 .

third lead. We neglect the Coulomb interaction in the grain. As we will see in the next section, the Coulomb interaction does not affect the main results we discuss here. In order to model the finite-size grain connected to a normal reservoir, we choose for convenience a finitesize wire characterized by its length l or equivalently by its level spacing $\Delta \sim \hbar v_F/l$. In fact, the precise shape of the finite-size grain is not important for our purpose as soon as it is characterized by a mean level spacing Δ separating peaks in the electronic density of states. We assume that the small quantum dot is weakly coupled to one or two adjacent leads (L and R). On the Hamiltonian level, we use the following tight-binding description, and for simplicity model the leads as one-dimensional wires (this is by no means restrictive): $H = H_L + H_R + H_0 +$ $H_{dot} + H_{tun}$ with

$$H_L = -t \sum_{j=1,s}^{\infty} (c_{j,s,L}^{\dagger} c_{j+1,s,L} + h.c.) - \mu_L n_{j,s,L}$$
 (1)

$$H_0 = -t \sum_{j=1,s}^{\infty} (c_{j,s,0}^{\dagger} c_{j+1,s,0} + h.c.) - \mu_0 n_{j,s,0}$$
(2)
+ $(t - t') \sum (c_{j,s,0}^{\dagger} c_{j+1,s,0} + h.c.)$

$$+(t-t')\sum_{s}(c_{l,s,0}^{\dagger}c_{l+1,s,0}+h.c.)$$

$$H_{dot} = \sum_{s} \epsilon_{d} n_{d,s} + U n_{d\uparrow} n_{d\downarrow}$$
 (3)

$$H_{tun} = \sum_{s} \sum_{\alpha=L,R,0} (t_{\alpha} c_{ds}^{\dagger} c_{1,s,\alpha} + h.c.).$$
 (4)

 H_R is obtained from H_L by changing $L \to R$. Here $c_{j,s,\alpha}$ destroys an electron of spin s at site j in lead $\alpha = 0, L, R; c_{d,s}$ destroys an electron with spin s in the dot, $n_{j,s,\alpha}=c_{j,s,\alpha}^{\dagger}c_{j,s,\alpha}$ and $n_{ds}=c_{ds}^{\dagger}c_{ds}$. The quantum dot is described by an Anderson impurity model, ϵ_d , U are respectively the energy level and the Coulomb repulsion energy in the dot. The tunneling amplitudes

between the dot and the left lead, right lead and grain are respectively denoted as t_L, t_R, t_0 (see Fig. 1). The tunneling amplitude amplitude between the grain and the third lead is denoted as t' (see Fig. 1). Finally tdenotes the tight binding amplitude for conduction electrons implying that the electronic bandwidth $\Lambda_0 = 4t$. Since we want to use the left and right leads just as transport probes, we assume in the rest of the paper that $t_L, t_R \ll t_0$.

We are particularly interested in the Kondo regime where $\langle n_d \rangle \sim 1$. In this regime, we can map $H_{tun} + H_{dot}$ to a Kondo Hamiltonian by help of a Schrieffer-Wolff transformation:

$$H_K = H_{tun} + H_{dot} = \sum_{\alpha, \beta = L, R, 0} J_{\alpha\beta} c_{1, s, \alpha}^{\dagger} \frac{\vec{\sigma}_{ss'}}{2} \cdot \vec{S} c_{1, s', \beta},$$
(5)

where $J_{\alpha\beta} = 2t_{\alpha}t_{\beta}(1/|\varepsilon_d| + 1/(\varepsilon_d + U))$. It is clear that $J_{00} \gg J_{0L}, J_{0R} \gg J_{LL}, J_{RR}, J_{LR}$. In Eq. (5), we have neglected direct potential scattering terms which do not renormalize and can be omitted in the low energy limit.

Kondo temperatures

The Kondo temperature is a crossover scale separating the high temperature perturbative regime from the low temperature one where the impurity is screened. There are many ways to define such scale. We choose the "perturbative scale" which is defined as the scale at which the second order corrections to the Kondo couplings become of the same order of the bare Kondo coupling. Note that all various definitions of Kondo scales differ by a constant multiplicative facor (see for example Ref. [8] for a comparison of the perturbative Kondo scale with the one coming from the Slave Boson Mean Field Theory).

The renormalization group equations relate the Kondo couplings defined at scales Λ_0 and Λ . They simply read:

$$J_{\alpha\beta}(\Lambda) \approx J_{\alpha\beta}(\Lambda_0)$$

$$+ \frac{1}{2} \sum_{\gamma} J_{\alpha\gamma}(\Lambda_0) J_{\gamma\beta}(\Lambda_0) \left[\int_{\Lambda}^{\Lambda_0} + \int_{\Lambda}^{-\Lambda} \right] \frac{\rho_{\gamma}(\omega)}{|\omega|} d\omega$$

where ρ_{γ} is the LDOS in lead γ seen by the quantum dot. When the density of states ρ_{γ} are uniform, the RG equations can be rewritten by introducing $\hat{\lambda}$, the matrix of the dimensionless Kondo couplings: $\lambda_{\alpha\beta} = \sqrt{\rho_{\alpha}\rho_{\beta}}J_{\alpha\beta}$. ^{9,14} The Kondo temperature T_K may be therefore defined as

$$\frac{1}{2} \left[\int_{T_K}^{\Lambda_0} + \int_{-\Lambda_0}^{-T_K} \frac{Tr(\hat{\lambda}(\omega))}{|\omega|} d\omega = 1. \right]$$
 (7)

Since $J_{00} \gg J_{LL}, J_{RR}$, the Kondo temperature essentially depends on the LDOS in the lead 0 and the Kondo temperature definition can be well approximated by

$$\frac{J_{00}}{2} \left[\int_{T_K}^{\Lambda_0} + \int_{-\Lambda_0}^{-T_K} \frac{\rho_0(\omega)}{|\omega|} d\omega = 1 \right]$$
 (8)

When the lead 0 becomes infinite (i.e. when t'=t), $\rho_0(\omega)=\rho_0=const$ and we recover $T_K=T_K^0$ the usual Kondo temperature where a constant density of states is assumed. It is worth noting that including the Coulomb interaction in the grain does not affect much the Kondo temperature. The grain Coulomb energy E_G slightly renormalizes J_{00} in the Schrieffer-Wolff transformation 12 since $E_G \ll U, |\epsilon_d|$.

The LDOS ρ_0 can be easily computed and corresponds in the limit $t'^2 \ll t^2$ to a sum of peaks at positions $\omega_n = -2t \cos k_{0,n} - \mu_0$ of width $\gamma_n \approx \frac{2(t')^2 \sin^3(k_{0,n})}{tl}$ where $k_{0,n} \approx \pi n/(l+1) + O(t'^2/t^2l)$ (Ref. 9). The LDOS ρ_0 is very well approximated by a sum of Lorentzian functions in the limit $t' \ll t$:

$$\pi \rho_0(\omega) \approx \frac{2}{l+1} \sum_n \sin^2(k_{0,n}) \frac{\gamma_n}{(\omega - \omega_n)^2 + \gamma_n^2}.$$
 (9)

This approximation is quite convenient in order to estimate the Kondo temperature T_K through (8). Note that the position of the resonance peaks may be controlled by the chemical potential μ_0 . Another possibility is to fix μ_0 and to add a small transverse magnetic field which modifies the orbital part of the electronic wave function in the grain and therefore shift the resonance peak positions. When the level spacing $\Delta_n \sim 2\pi t \sin k_{0,n}/l$ is much smaller than the Kondo temperature T_K^0 , no finite-size effects are expected. Indeed, the integral in (8) averages out over many peaks and the genuine Kondo temperature is $T_K \sim T_K^0$. On the other hand, when $T_K^0 \sim \Delta_n$, the Kondo temperature begins to depend on the fine structure of the LDOS ρ_0 and a careful calculation of the integral in (8) is required. Two cases may be distinguished: either ρ_0 is tuned such that a resonance ω_n sits at the Fermi energy $E_F = 0$ (labeled by the index R) or in a non resonant situation (labeled by the index NR).

In the former case, we can estimate

$$T_K^R = \frac{\gamma_n \Delta_n}{\sqrt{(\Delta_n^2 + \gamma_n^2) \exp\left(\frac{2}{J_{00}(\Delta_n)\rho_0^R(0)}\right) - \Delta_n^2}}$$

$$\approx \gamma_n \exp\left(-\frac{1}{J_{00}(\Delta_n)\rho_0^R(0)}\right) \approx \gamma_n \left(\frac{T_K^0}{\Delta_n}\right)^{\frac{\pi \gamma_n}{\Delta_n}} (10)$$

where we approximate the on-resonance LDOS at $E_F = 0$ by $\pi \rho_0^R(0) \approx 2 \sin^2 k_{0,n}/l\gamma_n$. In the latter case, we obtain,

$$T_K^{NR} = \Delta_n \exp\left(-\frac{\pi l \Delta_n^2}{16J_{00}(\Delta_n)\sin^2(k_{0,n})\gamma_n}\right)$$

$$\approx \Delta_n \exp\left(-\frac{1}{J_{00}(\Delta_n)\rho_0^{NR}(0)}\right) \approx \Delta_n \left(\frac{T_K^0}{\Delta_n}\right)^{\frac{\pi \Delta_n}{8\gamma_n}} \tag{11}$$

where $\pi \rho_0^{NR}(0) \approx 16\gamma_n \sin^2 k_{0,n}/l\Delta_n^2$. These two scales are very different when $t'^2 \ll t^2$. By controlling ρ_0 , we can control the Kondo temperature (only when $T_K \leq \Delta$). The main feature of such geometry is that the screening of the artificial spin impurity is essentially performed in the open finite-size wire corresponding to lead 0. Now we want to study what are the consequences of FSE on transport when one or two leads are weakly coupled to the dot. This is the purpose of the next section.

III. TRANSPORT SPECTROSCOPY OF A QUANTUM DOT COUPLED TO AN OPEN GRAIN

In this section, we consider a standard three-terminal geometry as depicted in Fig. 1 and analyze the conductance matrix of the system. There are several approaches we may combine to obtain such quantity for the whole temperature range. Nonetheless, before going into these details, let us analyze the dot density of states in presence of FSE in the grain.

A. Density of states

We have used the Slave Boson Mean Field Theory $(\operatorname{SBMFT})^{15}$ in order to calculate the dot density of states. This approximation describes qualitatively well the behavior of the Kondo impurity at low temperature $T \leq T_K$ when the impurity is screened. This method has been proved to be efficient to capture finite size effects in Ref. 9. The main advantage of the SBMFT relies on its ability to qualitatively reproduce the energy scales of the problem (here the Kondo temperature).

An interesting quantity to look at is the dot density of states $\rho_d(\omega)$. The density of states can be read from the differential conductance as follows: The current in the left electrode I_L reads:¹⁶

$$I_L = \frac{4e}{h} \Gamma_L \int_{-\infty}^{\infty} \left[2\pi f(\omega - \mu_L) \rho_d(\omega) - G_d^{<}(\omega) \right] d\omega, \quad (12)$$

where ρ_d is the dot LDOS, $G_d^<(\omega) = \int e^{i\omega t} \langle d^\dagger d(t) \rangle$ the lesser Green function for the dot and $\Gamma_{L/R} = \pi t_{L/R}^2 \rho_{L/R}$. The standard procedure is to get rid of the lesser dot Green function using the current conservation (here $I_L + I_R + I_0 = 0$). Nevertheless, such procedure is useful only when the leads density of states can be assumed constant on the typical scale we are interested in. This is not the case here because of the variations of $\Gamma_0(\omega) = \pi t_0^2 \rho_0(\omega)$. Nevertheless, one can make progress by assuming $\Gamma_{L/R} \ll \Gamma_0(\omega)$ such that for low bias the dot Green functions weakly depend on the chemical potential

in the left and right leads. Therefore

$$e\frac{dI}{d\mu_L} \approx \frac{2e^2}{h} 4\Gamma_L \int_{-\infty}^{\infty} \left(\frac{-df(\omega)}{d\omega}\right) \pi \rho_d(\omega + \mu_L) d\omega, \quad (13)$$

which allows an experimental access to $\rho_d(\mu_L)$. Note that a similar approximation is used for STM theory with magnetic adatoms¹⁷. We have plotted $\rho_d(\omega)$ in Fig. 2 for both the non-resonant case and the on-resonance case for three different values of ξ_K^0/l . We took the following parameters in units of t=1: $t_0=0.5,\ t_L=t_R=0.1$ (therefore $t_L^2<< t_0^2$), t'=0.5, and $l\sim 1000a$ (a the lattice constant) or equivalently $\Delta \sim 0.006$. The Kondo energy scale ξ_K^0 can be varied using the dot energy level ϵ_d which is controlled by the dot gate voltage. When $T_K^0 \gg \Delta$, no finite-size effect is to be expected. In this case, $\rho_d(\omega)$ mimics the density of state in the lead 0 but is shifted such that an off-resonance peak in the lead 0 corresponds to a dot resonance peak. The various peaks appearing in ρ_d are included in an envelope of width $O(T_K^0) \gg \Delta$ (which is a broader range than the figure 2 actually covers for $\xi_K^0/l = \Delta/T_K^0 \sim 0.15$). This can be simply understood from a non-interacting picture valid at T=0. The non-interacting dot Green function reads

$$G_{dd}(\omega) \approx \frac{1}{\omega - \epsilon_d - \delta\epsilon(\omega) + i\Gamma_0(\omega)},$$
 (14)

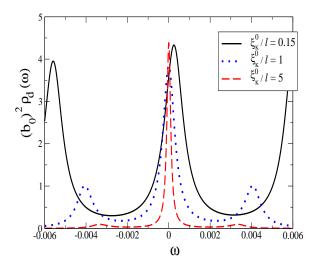
where $\delta\epsilon$ is the real part of the dot self-energy and Γ_0 its imaginary part. The minima's of Γ_0 thus correspond to the maxima of $-Im(G_{dd})$. We also note that the resonance peak is slightly shifted from $\omega=0$ in this limit. Therefore the 2-terminal conductance does not reach its unitary limit (i.e. its maximum non interacting value). This is due to the fact that we took $\epsilon_d=-0.68$ and we are not deep in the Kondo regime. Particle-hole symmetry is not completely restored in the low energy limit.

On the other hand, when $\xi_K^0 \gg l$ $(T_K^0 \ll \Delta)$, ρ_d changes drastically. The fine structure in the grain density of states no longer shows up in ρ_d . In the offresonance case, only the narrow Kondo peak of width $T_K^{NR} \ll T_K^0$ mainly subsists for $\xi_K^0/l = 5$ (upper panel of Fig. 2). We can also show that the position of the small peaks at $\omega \sim \pm \Delta/2$ for $\xi_K^0 \gg l$ are related to the resonance peaks in lead 0. We also note that the narrow peak is this time almost at $\omega = E_F = 0$ i.e. particle-hole symmetry is restored. In order to reach large value of ξ_K^0 , we took small values of ϵ_d such that we are deep in the Kondo regime where $n_D \sim 1$.

In the resonant case, the Kondo peak is split for $\xi_K^0 \gg l$. By approximating

$$\Gamma_0(\omega) \approx t_0^2 \sin^2(k_{0,n}) \frac{\gamma_n/l}{\omega^2 + \gamma_n^2},\tag{15}$$

at a resonance n, one can well understand the structure of ρ_d with the SBMFT. When $T_K^R \gg \gamma_n$, one can show that the peak splitting is of order $\sim 2\sqrt{\gamma_n T_K^R}$ and the peaks



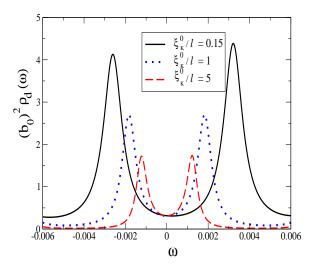


FIG. 2: Dot density of states $\rho_d(\omega)$ for both the non resonant case (upper panel) and the resonant case (lower panel). We took $\Delta \sim 0.006$ and plot ρ_d for $\xi_K^0/l = \Delta/T_K^0 \sim 0.15$, $\xi_K^0/l = \Delta/T_K^0 \sim 1$ and $\xi_K^0/l = \Delta/T_K^0 \sim 5$. Note that ρ_d has been scaled by b_0^2 , the slave boson parameter for an easy comparison between both cases.

width is of order γ_n . We have shown that the dot density of states $\rho_d(\omega)$ has interesting features at T=0 where FSE can be clearly identified. Let us summarize them: First, when $T_K^0 \gg \Delta$, ρ_d has several peaks embedded in an enveloppe of width $\sim T_K^0$ associated with the grain one-particle levels and one or two Kondo peaks in the opposite limit $T_K^0 \ll \Delta$. Second, the position of the peaks and their widths depend on the grain being on- or off- resonance.

B. Finite temperature conductances

In this section, we analyze the conductance matrix using various methods. We start with the low temperature limit.

1. Low temperature regime

At T=0, it is straightforward to show, using for example the scattering formalism,¹⁸ that the conductance matrix $G_{\alpha,\beta}^U$ is simply given by

$$G_{\alpha,\beta}^{U} = \frac{2e^2}{h} \frac{4\Gamma_{\alpha}\Gamma_{\beta}}{(\Gamma_L + \Gamma_0 + \Gamma_R)^2}$$
 (16)

where $\Gamma_{\alpha} = \pi t_{\alpha}^2 \rho_{\alpha}(0)$, and $\alpha = l, 0, r$. Since the SBMFT aims at replacing the initial Anderson Hamiltonian by a non-interacting one, one may easily access the conductance by directly applying the Landauer formula or equivalently by using

$$G_{\alpha\beta} = \frac{2e^2}{h} \int d\omega \left(\frac{-\partial f}{\partial \omega}\right) \frac{4\Gamma_{\alpha}(\omega)\Gamma_{\beta}(\omega)}{\left(\sum_{\alpha} \Gamma_{\alpha}(\omega)\right)} Im(-G_{dd}^r)(\omega). \tag{17}$$

We have plotted in Fig. 3, the conductance between the left lead L and the lead 0 for both the on- and off-resonance cases. Note that in absence of Coulomb energy the density of states in the grain can be controlled by the chemical potential μ_0 .

The three upper curves correspond to the non-resonant cases whereas the three lower curves correspond to the resonant cases. In the standard case corresponding to $\xi_K^0 \ll l \ (T_K^0 \gg \Delta)$, the conductance qualitatively follows the non interacting limit except at high temperature (here $T \gg T_K^0 \sim 0.012$) where the conductance falls down in the Kondo regime. The conductances corresponding to $\xi_K^0 \gg l$ are characterized by an abrupt increase of the conductance at their respective Kondo temperatures T_K^R and T_K^{NR} which are lower than T_K^0 .

The main signature of finite size effects occurs at an intermediate temperature $T_K^0 > T \sim 0.001 > T_K^R, T_K^{NR}$. In this temperature range, the conductor is already a fraction of its unitary limit value when $\xi_K^0/l \sim 0.15 \ll 1$ whereas it is still very low when $\xi_K^0/l \sim 5 > 1$. In this temperature range, significant deviations from the non-interacting limit are obtained for $\xi_K^0/l \sim 5$ contrary to the case where $\xi_K^0/l \sim 0.15$. We also notice that the conductance for the on-resonance case is non monotonous. This is analyzed and confirmed analytically further using Nozières' Fermi liquid approach. As anticipated before from the spectroscopic analysis, the unitary limit is not fully reached for the NR case at $\xi_K^0 < l$. It corresponds to the fact that particle-hole symmetry is not fully restored at low energy in the SBMFT approximation. Because of the large value of T_K^0 we took (in order to satisfy $\xi_K^0 < l$) the dot occupation n_d starts slightly deviating from 1

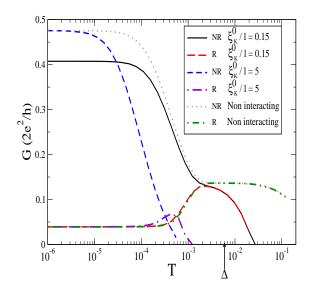


FIG. 3: Conductance between lead L and lead 0 (in units of $2e^2/h$) as a function of temperature when the density of states in the grain is tuned on resonance (curves denoted as R) and out of resonance (curves denoted as NR). Three cases were considered: $\xi_K^0/l \sim 0.15$ (lower plain style and upper long dashed curves), $\xi_K^0/l \sim 5$ (lower dashed and upper dot-dashed curves). We compare these curve to the non interacting case corresponding to $\epsilon_d = U = 0$ (dotted lines). The level spacing Δ is indicated by an arrow on the x axis.

which explains why the unitary limit is not fully restored at low T.

Similar results can be obviously obtained by analyzing the conductance G_{LR} despite the fact that the amplitude of G_{LR} will be in general smaller than G_{L0} by a factor $\sim \sqrt{\Gamma_L \Gamma_R}/\Gamma_0$. In fact, the conductance G_{LR} can be made significantly larger by adjusting the chemical potential μ_0 such that $\langle I_0 \rangle = 0$. The grain or OFSW density of states ρ_0 can then be controlled by a small orbital magnetic field or by gating the grain. In the linear response regime, this can always be realized provided $\mu_0 = (G_{L0}\mu_L + G_{R0}\mu_R)/(G_{L0} + G_{R0})$. In this sense, the geometry becomes a two-terminal one. More generally, one can also show using a general non-equilibrium approach following Ref. [16] that there is a unique value of μ_0 such that $I_0 = 0$. Note that for a symmetric device (between left and right) and $\mu_L = -\mu_R = eV/2$, symmetry considerations imply $I_0 = 0$ for $\mu_0 = 0$. For voltages smaller than the smallest energy scale of the problem (T_K^0) or γ_n), we can safely use the simplified Meir-Wingreen formula. 16 This "two-terminal like" conductance then

$$G_{LR}^{(2)} = -\frac{2e^2}{h} \int d\omega \left(\frac{-\partial f}{\partial \omega}\right) \Gamma_{LR}(\omega) Im(G_{dd}^r)(\omega), \quad (18)$$

where $\Gamma_{LR} = \frac{4\Gamma_L\Gamma_R}{(\Gamma_L + \Gamma_R)}$. We have added the index (2)

to distinguish it from the 3-terminal conductance. Contrary to Eq. (17), this equation is exact and does not rely on a mean-field approximation. Under the approximation $t_L, t_R \ll t_0$, one may relate the exact dot Green's function to the on shell T-matrix associated with the scattering problem of an electron in lead 0 by the dot: $T(\varepsilon) \approx t_0^2 G_{dd}^T$ such that:

$$G_{LR}^{(2)} \approx \frac{2e^2}{h} \int d\omega \left(\frac{-\partial f}{\partial \omega}\right) \frac{\Gamma_{LR}(\omega)}{\Gamma_0(\omega)} \left(-\pi \rho_0(\omega) Im(T(\omega))\right).$$
 (19)

We see that by adjusting μ_0 such that $I_0 = 0$, the amplitude of $G_{LR}^{(2)}$ can be made of the order of G_{L0} . We have checked that $G_{LR}^{(2)}(T)$ has the same features as $G_{L0}(T)$ as it should be

2. Nozières' Fermi liquid approach

Equation (19) may be good starting point to analyze the finite-temperature Fermi liquid correction to the unitary limit. In the Fermi liquid approach the on-shell T-matrix is expanded at low T as a function of ω and T. Here new energy scales are introduced and this expansion must be done with care. Let us start with the off-resonance case where at low energy $\rho_0(\omega) \sim const$. In this case, one can safely use the usual T- matrix expansion T-

$$-\pi \rho_0(\omega) Im(T(\omega)) \approx 1 - \frac{3\omega^2 + \pi^2 T^2}{2(T_K^{NR})^2}.$$
 (20)

We then recover the standard Fermi liquid corrections

$$G_{LR}^{(2)}(T) \approx G_U^{(2)} \left(1 - \pi^2 \left(\frac{T}{T_K^{NR}} \right)^2 \right) ,$$
 (21)

where $G_U^{(2)} = 4\Gamma_L\Gamma_R/(\Gamma_0(\Gamma_L\Gamma_R))$. Therefore the off-resonance conductance always decreases at low temperature as depicted in Fig. 3. However the situation is more subtle when ρ_0 is on a resonance.

In order to perform an expansion of the on-shell T-matrix analogous to Eq. (20), one should first pay attention to the new energy scale γ_n . When $\gamma_n \gg T$, one may reproduce the same expansion as in [19]. Eq. (20) remains qualitatively valid provided we replace T_K^{NR} by T_K^R . Reporting together both Eqs (20) and (15) in Eq. (19) and performing a Sommerfeld expansion, we find:

$$G_{LR}^{(2)}(T) \approx G_U^{(2)} \left(1 + \pi^2 T^2 \left(\frac{2}{3(\gamma_n)^2} - \frac{1}{(T_K^R)^2} \right) \right), \quad (22)$$

We find two opposite contributions to the conductance at low temperature: the usual negative one coming from the T-matrix expansion and a second positive one depending on the resonance width. The latter has nothing to do with interactions and is simply an effect of quantum interference analogous to the Fano resonance recently seen

in quantum dot experiments.²⁰ For a narrow resonance satisfying $\gamma_n \ll T_K$, the conductance increases at low $T \ll \gamma_n$. On the other hand, for a broad resonance, we recover the usual situation. Note that this value suggests an intermediate value of γ_n where the T^2 corrections cancel. For the 3-terminal conductance matrix, we can perform a similar analysis starting from Eq. (17) which can be regarded as a good approximation of the conductance in Nozières' Fermi liquid regime and obtain almost similar conclusions.

3. High temperature

At temperature higher than the Kondo temperature $T\gg T_K^0$, perturbation theory in the Kondo couplings can a priori be used. We therefore start directly from the Kondo Hamiltonian written in Eq. (5) In order to compute the conductance between the left lead and lead 0, we use the renormalized perturbation theory. At lowest order, the conductance reads:

$$G_{L0} = \frac{2e^2}{h} \frac{3\pi^2}{4} J_{L0}^2 \int d\omega \left(\frac{-\partial f}{\partial \omega}\right) \rho_L \rho_0(\omega)$$
 (23)

When $T_K^0 \gg \Delta$, one may replace ρ_0 by its average value and J_{L0} by its renormalized coupling and the high temperature conductance takes its standard scaling form²¹

$$G_{L0} = G_{L0}^{U} \frac{3\pi^{2}/16}{\ln^{2}(T/T_{K}^{0})}, T \gg T_{K}^{0} \gg \Delta$$
 (24)
= $G_{L0}^{U} f(\frac{T}{T_{K}^{0}}),$

where f(x) is a universal scaling function such that $f(x\gg 1)\approx \frac{3\pi^2}{16\ln^2 x}$ and G_{L0}^U has been defined in Eq. (16).

Let us consider the more interesting situation where $T_K^0 < \Delta$. Suppose the bandwidth is reduced from $\pm \Lambda_0$ (where $\lambda_0 = 4t$ here) to $\pm \Lambda$, the renormalization of the Kondo coupling is well approximated by:

$$J_{L0}(\Lambda) = J_{L0}(\Lambda_0) + \frac{1}{2} J_{L0} J_{00} \left[\int_{-\Lambda_0}^{-\Lambda} + \int_{\Lambda}^{\Lambda_0} d\omega \frac{\rho_0(\omega)}{|\omega|} \right]. \tag{25}$$

When $\Lambda \gg \Delta$, the integral in Eq. (25) averages over many peaks in the LDOS ρ_0 and we obtain the usual result for the Kondo model with a constant density of states. It implies that in the limit $T \gg \Delta$, the result in Eq. (24) remains valid independently of $\rho_0(\omega)$ being tuned on or off a resonance.

On the other hand, when $\Lambda \ll \Delta$, the integral in Eq. (25) becomes strongly dependent on ρ_0 . When ρ_0 is tuned off a resonance and for $\Delta \gg T \gg T_K^{NR}$, the result in Eq. (24) remains approximately valid provided we replace T_K^0 by T_K^{NR} defined in Eq. (11) such that $G_{L0}(T) = G_U f(T/T_K^{NR})$. When ρ_0 is tuned on

a resonance that we assume for convenience to be at $\omega = E_F = 0$, only the variation of the LDOS in the vicinity of E_F matters such that we further approximate the LDOS using Eq. (15). The renormalization group equation in Eq. (25) may be integrated in two steps, first between Λ_0 and Δ where the variations of the LDOS average out and then between Δ and Λ where

$$J_{L0}(\Lambda) \approx J_{L0}(\Delta) + \frac{1}{2} J_{L0}(\Delta) J_{00}(\Delta) \rho_0(0) \ln\left(1 + \frac{\gamma_n^2}{\Lambda^2}\right),$$

where we assumed $\gamma_n \ll \Delta$. At $\Lambda \gg \Delta$, the Kondo couplings renormalize following the RG equations given in Eq. (6). At the scale Δ , the Kondo couplings have been weakly renormalized and one may use the RG equations given in Eq. (6) that imply $J_{L0}(\Delta) \approx \frac{t_1 t_0}{t_0^2} J_{00}(\Delta)$. Using this approximation and the definition of the onresonance Kondo temperature in Eq. (10), the conductance at $\Delta > T \gg \gamma_n$ reads

$$G_{L0} \approx G_U \frac{3\pi^2}{16} \frac{1}{\ln^2(1 + \frac{\gamma_n^2}{(T_K^R)^2})} \frac{\gamma_n}{T} \left(1 + \frac{\ln(1 + \frac{\gamma_n^2}{T^2})}{\ln(1 + \frac{\gamma_n^2}{(T_K^R)^2})} \right)^2. \tag{27}$$

The high temperature on-resonance conductance takes a more complicated form that the one given in Eq. (24). Notice nonetheless that the multiplicative factor $(1+\cdots)$ appearing in Eq. (27) is O(1) since $T\gg T_K^R=O(\gamma)$.

C. Scaling analysis

From the low and high temperature analysis, we have seen that the conductance matrix elements cannot be in general simply written as a simple universal scaling function of T/T_K^0 both at high and low temperature. This is particularly striking when ρ_0 was tuned on a resonance characterized by the width γ_n (see Eqs (22) and (27)).

In general, one would expect

$$G_{\alpha\beta} = G_{\alpha\beta}^{U} g(\frac{T}{T_{K}^{0}}, \frac{\Delta}{T_{K}^{0}}, \frac{\gamma_{n}}{T}). \tag{28}$$

In the high temperature regime, this scaling function takes a simple form and simply reads $f(T/T_K)$ where T_K either equates T_K^0 when $\Delta \ll T_K^0$ or can be simply expressed as a function of T_K^0, Δ, γ_n (see eqs (10) and (11)). Nevertheless, at intermediate or low temperature, the conductance in Eq. (28) takes a complicated scaling form which depends on γ_n/T and is in this sense non universal. It depends on the geometric details of the sample, at least for bare Kondo temperature T_K^0 of the order or smaller than the level spacing Δ . A similar conclusion has been reached by Kaul et al. 22 by assuming a chaotic large quantum grain and explicitly calculating the fluctuations and deviations from the universal behavior taking into account mesoscopic fluctuations.

D. Finite grain Coulomb energy

In this section, we discuss whether a finite grain Coulomb energy modifies or not the results presented in this work. As we already mentioned in section 2, the Kondo coupling J_{00} , is almost not affected by the grain Coulomb energy E_G (since $E_G \ll U$) and therefore the Kondo temperature remains almost unchanged. As shown in 12,23,24, a small energy scale E_G changes the renormalization group equation in Eq. (6). The offdiagonal couplings $J_{0L}(\Lambda)$, $J_{0R}(\Lambda)$ tend to 0 for $\Lambda \ll E_G$. At energy $\Lambda \ll E_G$ the problem therefore reduces to an anisotropic 2-channel Kondo problem. The strongly coupled channel is the grain 0, the weakly coupled one is the even combination of the conduction electron in the left/right leads. At very low energy, the fixed point of the anisotropic 2-channel Kondo model is a Fermi liquid. It is characterized by the strongly coupled lead (here the grain) screening the impurity whereas the weakly coupled one completely decouples from the impurity. The dot density of states depicted in Fig. 2 remains therefore almost unaffected. The on and off-resonance Kondo temperatures T_K^R and T_K^{NR} given in Eqs (10) and (11) remain valid too. The problem is to read the dot LDOS with the weakly coupled leads since they decouple at T=0. Nevertheless, for a typical experiment done at low temperature T, such a decoupling is not complete and the dot density of state should be still accessible using the weakly coupled leads but with a very small amplitude.

In this paper we analyze the situation in which a grain or a finite size wire is also used as a third terminal. In some situations, like the theoretical one presented in [12], no terminal lead is attached to the grain and the geometry is a genuine 2-terminal one. Taking into account both a finite level spacing and a finite grain Coulomb energy is quite involved (with several different regimes) and goes beyond the scope of the present paper. A step into this direction was recently achieved in Ref. [25].

IV. CONCLUSIONS

In this paper, we have studied a geometry in which a small quantum dot in the Kondo regime is strongly coupled to a large open quantum dot or open finite size wire and weakly coupled to other normal leads which are simply used as transport probes. The artificial impurity is mainly screened is the large quantum dot. Such a geometry thus allows to probe the dot spectroscopic properties without perturbing it. We have shown combining several techniques how finite size effects show up in the dot density of states and in the the conductance matrix. We hope the predictions presented here are robust enough to be checked experimentally.

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